

**Maryland Pharmacy Program**  
**HEPATITIS C PROTEASE INHIBITORS PRIOR-AUTHORIZATION**

Phone: 410-767-1755 or 1-800-492-5231-Option 3

Fax form to: 410-333-5398

Incomplete forms will be returned-A copy of the patient medical history with required lab test results must accompany this request.

**Section I. Patient Information**

Recipient: \_\_\_\_\_ MA#: \_\_\_\_\_ Date of Birth: \_\_\_\_/\_\_\_\_/\_\_\_\_  
Phone #: (\_\_\_\_) \_\_\_\_\_ - \_\_\_\_\_ Patient location: \_\_\_\_ Home \_\_\_\_ Hospital \_\_\_\_ Clinic \_\_\_\_ Office  
Address: \_\_\_\_\_

**Section II- Drug/Drug Combination is used for an FDA-approved indication**

List Hepatitis C triple drug regimen with strengths/dosages:

☐ Victrelis: \_\_\_\_\_ ☐ Incivek: \_\_\_\_\_  
☐ Pegasys: \_\_\_\_\_ ☐ PegIntron: \_\_\_\_\_  
☐ Ribavirin: \_\_\_\_\_ (Only the generic 200mg ribavirin tab/cap will be approved)

Diagnoses: \_\_\_\_\_ Genotype: ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5 ☐ 6

☐ Acute Hep C ☐ Chronic Hep C ☐ Other \_\_\_\_\_ Body Weight: \_\_\_\_\_ kg

Baseline HCV RNA level: \_\_\_\_\_ Date measured: \_\_\_\_\_

HCV RNA Level at TW 4: \_\_\_\_\_ Date measured: \_\_\_\_\_

at TW 8: \_\_\_\_\_ Date measured: \_\_\_\_\_

at TW 12: \_\_\_\_\_ Date measured: \_\_\_\_\_

at TW 24: \_\_\_\_\_ Date measured: \_\_\_\_\_

at TW \_\_: \_\_\_\_\_ Date measured: \_\_\_\_\_

Liver enzyme levels: ALT/AST: \_\_\_\_\_ Date measured: \_\_\_\_\_

Does patient have HIV/HCV co-infection? ☐ Yes ☐ No

Has a liver biopsy been performed? ☐ Yes ☐ No If yes, date: \_\_\_\_\_ Provide a copy of biopsy results.

Is this for New Therapy? ☐ New ☐ Relapser ☐ Partial responder ☐ Non-responder

Does the patient have a history of any of the following:

\_\_\_ Anemia; \_\_\_ Major uncontrolled depressive illness; \_\_\_ Alcohol or drug abuse; \_\_\_ Untreated thyroid d/s;  
\_\_\_ Autoimmune hepatitis or other autoimmune conditions; \_\_\_ Pregnant (ribavirin is category X); \_\_\_ Thrombocytopenia;  
\_\_\_ Severe concurrent medical d/s (ie. AIDS, cancer, significant CAD); \_\_\_ Decompensated liver d/s;  
\_\_\_ Hemoglobinopathies (i.e. sickle cell, thalassemia); \_\_\_ on didanosine; \_\_\_ On concomitant interacting drugs;  
\_\_\_ Unstable CVD; \_\_\_ Solid organ transplant (renal, heart, or lung); Other: \_\_\_\_\_

Is patient a good candidate for the triple drug regimen in terms of compliance to this complex regimen? ☐ Yes; ☐ No

Any prior side-effects/adverse effects from prior therapy, i.e. peg-interferon alfa and/or ribavirin? ☐ Yes; ☐ No

List prior failed drug regimens for Hep C and reasons for drug failure: \_\_\_\_\_

Provide reasons for selecting one Hep C protease inhibitor over the other: \_\_\_\_\_

**Section III- Drug/Drug Combination is used off-label**

Off-label indications or off-label dosage: \_\_\_\_\_

Is drug(s) used as part of a clinical study or trial? ☐ No ☐ Yes- If yes, specify sponsoring organization/drug manufacturer: \_\_\_\_\_

Specify purpose of study: \_\_\_\_\_

List references and reasons supporting off-label uses: \_\_\_\_\_

**Note:** Off-label indications or off-label dosages may be approved if medically necessary, safe, appropriate, and documented in/supported by one of the three official compendia (AHFS Drug Information, Micromedex Drugdex, and US Pharmacopeia). The Program takes into consideration the Drugdex's ratings related to the strength of recommendation and the evidence on drug safety and efficacy.

I certify that the information provided is accurate. Supporting documentation is available for State audits.

\_\_\_\_\_, M.D. Prescriber's Name: \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_  
(Prescriber's signature). Tel# (\_\_\_\_) \_\_\_\_\_ - \_\_\_\_\_ Fax# (\_\_\_\_) \_\_\_\_\_ - \_\_\_\_\_  
License #: \_\_\_\_\_ DEA #: \_\_\_\_\_ Specialty : \_\_\_\_\_  
Address: \_\_\_\_\_

